NITROGEN CONTAINING COMPOUND

1.0 Amines

Amines are derivatives of ammonia in which one or more hydrogen atoms are replaced by alkyl group(s).

Amines are classified as primary, secondary and tertiary depending on the number of alkyl groups attached to nitrogen atom.

$$NH_{3} \xrightarrow{+R} \longrightarrow RNH_{2} \xrightarrow{+R} \xrightarrow{+R} R_{2}NH \xrightarrow{-H} R_{3}N$$

$$p-amine \qquad sec. amine \qquad tertiary amine$$

$$General formula (C_{n}H_{2n+3}N)$$

$$\downarrow \qquad \qquad \downarrow$$

$$1^{\circ} amine \qquad 2^{\circ} amine \qquad 3^{\circ} amine$$

2.0 General methods of preparation

- (1) Ammonolysis of alkyl halides and alcohols:
 - (a) From Ammonolysis of alkyl halides [Hofmann's ammonolysis]: When an aqueous solution of ammonia is heated with alkyl halide all the three types of amines and quaternary ammonium salt are formed

$$R - X \xrightarrow{NH_3} R - NH_2 \xrightarrow{R-X} R_2NH \xrightarrow{R-X} R_3N \xrightarrow{R-X} R_4NX$$
(Quaternary ammonium salt)

If ammonia is taken in excess, 1° amine is the main product.

(b) Ammonolysis of alcohols : When ROH and NH_3 are passed over Al_2O_3 or ThO_2 at 350° C all the three types of amines are formed.

$$\text{R--OH} \xrightarrow{\text{NH}_3 \atop \text{Al}_2\text{O}_3 / 400^\circ\text{C}} \text{R--NH}_2 \xrightarrow{\text{R-OH} \atop \text{Al}_2\text{O}_3} \text{R}_2\text{NH} \xrightarrow{\text{R-OH} \atop \text{Al}_2\text{O}_3} \text{R}_3\text{N}$$

- Quaternary ammonium hydroxide is not formed.
- If excess of ammonia is used, then main product will be primary amine.
- (2) By reduction:

(a) With RCONH₂:
$$RCONH_2$$
 $\xrightarrow{LiAlH_4}$ RCH_2NH_2

(b) With RCN:
$$RCN + 4[H] \xrightarrow{Na/C_2H_5OH} RCH_2NH_2$$

This reaction (b) is called mendius reaction.

The reduction of alkyl isocynides gives secondary amines.

(c) With Oximes:
$$R-CH=N-OH+4[H]$$
 $\xrightarrow{LiAlH_4 \text{ or}} RCH_2-NH_2 + H_2OH_2$

(d) With RNO₂: RNO₂ + 6[H]
$$\xrightarrow{\text{Sn/HCl}}$$
 RNH₂ + 2H₂O

Sn/HCl is used in laboratory preparation

- (3) By hydrolysis of:
 - (a) R—NC: Alkyl isocyanide undergoes hydrolysis with mineral acid and forms alkyl amine.

$$R-NC + 2H_2O \xrightarrow{HCl} RNH_2 + HCOOH$$



(b) RNCO: Alkyl isocyanate undergoes hydrolysis on heating with KOH and forms alkyl amine.

(4) By Hofmann's bromamide reaction (Hofmann's Hypobromite reaction) : This is a general method for the conversion of alkanamides into primary amines having one less carbon.

O II
$$R-C-NH_2 + Br_2 + 4KOH \longrightarrow R-NH_2 + K_2CO_3 + 2KBr + 2H_2O$$

(5) From Grignard reagent: Alkyl magnesium iodide reacts with chloramine to yield alkyl amine.

$$R\!\!-\!\!Mg\!-\!\!I \; + \; Cl\!-\!\!NH_{_{\!2}} \qquad \longrightarrow \qquad R\!\!-\!\!\!NH_{_{\!2}} \; + \; Mg \, \Big\langle \begin{matrix} I \\ Cl \end{matrix}$$

(6) Gabriel phthalimide synthesis : Phthalimide is first treated with KOH to obtain potassium phthalimide which is then treated with alkyl iodide. Then alkyl phthalimide on hydrolysis yields alkylamine. This method is used in the formation of pure aliphatic primary amines.

$$R-NH_{2} + COOH COOH COOH COONR$$

Phthalic acid

(7) Curtius reaction:

(8) Schmidt reaction: In presence of conc. H_2SO_4 alkanoic acid reacts with hydrazoic acid (N_3H) followed by hydrolysis to yield alkylamine.

$$R$$
— $COOH + N3H $\xrightarrow{(1) N_3H, H_2SO_4}$ R — $NH_2 + N_2 + CO_2$$

GOLDEN KEY POINTS

• Separation of 1°, 2° and 3° amines

Mixture of 1° , 2° , 3° amine can be separated by following methods.



^{*} Aniline is not formed by this reaction.

- **(i) Fractional distillation :** The mixture of amines may be separated by fractional distillation because their boiling points are quite different. It is used in industry.
- **(ii) Hinsberg method :** In this method mixture of amines is seperated by using benzene sulphonyl chloride (Hinsberg's reagent).

$$C_6H_5SO_2Cl + 1^\circ$$
 amine \longrightarrow Product \xrightarrow{KOH} Soluble (ppt.)

 $C_6H_5SO_2Cl + 2^\circ$ amine \longrightarrow Product \xrightarrow{KOH} insoluble (ppt.)

3° amine does not react with benzene sulphonyl chloride. (No ppt. formed)

(iii) Hofmann method: In this method mixture of amines is separated by using ethyl oxalate.

 1° amine + ethyl oxalate \longrightarrow solid product

 2° amine + ethyl oxalate \longrightarrow liquid product

 3° amine + ethyl oxalate \longrightarrow No reaction

3.0 PHYSICAL PROPERTIES

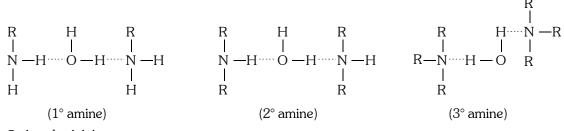
- (i) CH_3NH_2 is gas and $C_2H_5NH_2$ is a volatile liquid.
- (ii) Higher amines have fishy smell.
- (iii) H-Bonding (weaker as compared to H—O—H).

In 3° amine (due to absence of H-atom) H-bonding is not possible.

(iv) **Boiling point:** Due to weak intermolecular H-bonding the B.P. of 1° and 2° amines are lower than those of alcohols of comparable molecular weight. The boiling point of 3° amines which form no H-bonds are near to those of alkanes of comparable molecular weight.

Boiling point α molecular weight

(v) **Solubility:** Low molecular weight amines are soluble in water. The water solubility of amines decreases with increasing size of alkyl group.



Order of solubility \longrightarrow p- amine > s- amine > t- amine



4.0 Chemical properties

- (i) Basic character of amines is due to the presence of lone pair of electrons on the N atom.
- (ii) Basic strength depends on electron donating tendency.
- (iii) Order of basic character in aqueous solution : $(CH_3)_2 NH > CH_3 NH_2 > (CH_3)_3 N > NH_3$ $(C_2H_5)_2 NH > (C_2H_5)_3 N > C_2H_5 NH_2 > NH_3$

GOLDEN KEY POINTS

- Tertiary amine is less basic then secondary due to following reasons:
 - (i) **Steric hindrance :** In tertiary amines (R₃N) , three alkyl groups attached to N are bulkier and as such exert steric hindrance.
 - (ii) Decrease in hydration:

In tertiary amine

In secondary amine

$$R_3$$
N $-H$ ---O $\begin{pmatrix} H \\ H \end{pmatrix}$

 R_2 H O H

Protonated t-amine can form H-bonding with water molecule only at one point [less stable]

Protonated s—amine can form H—bonding with water molecules at two points (more stable)

Conjugate acid of 3° amine are less stable as compare to 2° amine due to low hydration so less basic.

- The basic strength of aniline is less than aliphatic amines as the lone pair of electron present on N- atom interact with the delocalized π orbital of benzene ring. Hence it is less available for protonation on N- atom.
 - (1) Reactions showing basic nature:

It reacts with acids to form salts.

$$RNH_2 + HCl \longrightarrow [RNH_3] \stackrel{\Theta}{Cl} \stackrel{\Delta}{\longrightarrow} R-Cl + NH_3$$

Alkyl ammonium chloride

(Acidic salt)

$$2RNH_2 \xrightarrow{H_2SO_4} (RNH_3)_2 SO_4^{-2}$$
 Alkyl ammonium sulphate

(2) Reaction with alkyl halides: Alkyl amine reacts with alkyl halides and form sec., ter. amines and quaternary ammonium salt.

$$RNH_2 + R-X \xrightarrow{-HX} R_2NH \xrightarrow{RX} R_3N \xrightarrow{RX} R_4NX$$

(3) Acetylation: Acetylation takes place when alkyl amine combines with acetyl chloride or acetic anhydride.

$$RNH_2 + CICOCH_3 \longrightarrow RNHCOCH_3 + HCI$$
 $(N -alkyl acetamide)$
 $RNH_2 + (CH_3CO)_2O \longrightarrow RNHCOCH_3 + CH_3COOH$
 $(N -alkyl acetamide)$



(4) Benzoylation (Schotten Baumann reaction):

$$\begin{array}{c} \text{COCl} & \text{CONHR} \\ \hline \bigcirc & + \text{ H-NH-R} & \xrightarrow{\text{NaOH}} & \hline \bigcirc & + \text{ HCl} \\ \end{array}$$

Benzoylchloride

N-alkyl benzamide

(5) Acidic nature: Amines are very weak acids only 1° and 2° amines show acidic nature with active metals.

N- alkyl sodamide

(6) Reaction with Tilden reagent:

$$R-NH_9 + NOCl \longrightarrow RCl + N_9 + H_9O$$

(7) Reaction with aldehydes:

$$H$$
 $R-NH_2 + O = C - R \xrightarrow{-H_2O} RCH = NR \text{ (Schiff's base)}$

(8) Carbylamine Reaction (Isocyanide test): When alkyl amine heated with chloroform and alc. KOH alkyl isocyanide is formed which has very bad smell.

This test is also given by aniline . This is a test for p-amines.

$$R-NH_2 + CHCl_3 + 3 KOH \longrightarrow R-N \cong C + 3KCl + 3H_2O$$

(9) Hofmann's mustard oil test: When alkyl amine is heated with carbon disulphide and mercuric chloride alkyl isothiocyanate is formed which has smell like mustard oil.

Alkyl isothiocyanate

$$\begin{array}{ccc} & & & & & \\ & & & & \\ R_3N & + & C = S & & \longrightarrow & & \text{No reaction.} \end{array}$$

(10) Reaction with HNO₂:

(a) Primary amines react with nitrous acid to produce nitrogen gas [seen as bubbles]

$$R-NH_2 + HONO \longrightarrow R-OH + N_2 \uparrow + H_2O$$

$$CH_3 NH_2 + HNO_2 \longrightarrow CH_3 - O - CH_3$$

(b)
$$R_2 NH + HONO \longrightarrow R_2 N-NO + H_2O$$

Dialkylnitroso amine (Yellow oily layer)

(c)
$$R_3N + HONO \longrightarrow R_3N HNO_2^{\Theta}$$
 Trialkyl ammonium nitrite (Soluble in water)



Points to Remember:

- (i) Nitrosoamines are carcinogens (Cancer causing agents)
- (ii) Amines can have chiral N-atom but cannot be resolved into enantiomeric forms because of rapid inversion of one enantiomeric form into the other.
- (iii) The Schiff's bases formed by reaction of 1° -amines and aldehyde/ketones are also called anils.
- (iv) The mixture of 1° , 2° , 3° amines can be distinguished by Hofmann's test or Hinsberg's reagent or nitrous acid test.

5.0 ANILINE (C₆H₅NH₉)

5.1 General Methods of Preparation

(1) Lab method:

$$C_6H_5NO_2 \xrightarrow{Sn+HCl} C_6H_5NH_2 + H_2O_1$$

(2) Industrial method:

$$C_6H_5$$
— NO_2 — $Fe+HCl$ C_6H_5 — NH_2 + H_2O

(3) From Phenol:

$$C_6H_5OH + NH_3 \xrightarrow{ZnCl_2} C_6H_5NH_2 + H_2O$$

(4) **From benzamide** (Hofmann's bromamide reaction):

$$C_6H_5CONH_2 + Br_2 + 4KOH \longrightarrow C_6H_5NH_2 + K_2CO_3 + 2KBr + 2H_2O$$

(5) **From benzoic acid** (Schmidt reaction):

(6) From Grignard reagent :

$$C_6H_5MgBr + ClNH_2 \longrightarrow C_6H_5NH_2 + Mg < Cl$$

(7) From phenyl isocyanide:

$$C_6H_5N \Longrightarrow C + 2H_2O \longrightarrow C_6H_5NH_2 + HCOOH$$

5.2 Physical Properties

- (i) Fresh, aniline is a colourless oily liquid. On standing the colour becomes dark brown due to action of air and light.
- (ii) It's B.P. is 183°C.
- (iii) It is heavier than water.
- (iv) It has characteristic unpleasent odour. It is toxic in nature.

Reactions due to -NH, group

(1) **Basic nature**: Aniline is weak base but it forms salt with strong acids. It accepts a proton.

$$C_6H_5\ddot{N}H_2 + H^+ \longrightarrow C_6H_5\ddot{N}H_3$$
 Anilinium ion

 $C_6H_5NH_2 + HCl \longrightarrow C_6H_5NH_3Cl$ Anilinium hydrochloride

 $2C_6H_5NH_2 + H_2SO_4 \longrightarrow (C_6H_5\ddot{N}H_3)_2SO_4$ Anilinium sulphate

 $2C_6H_5NH_2 + H_2PtCl_6 \longrightarrow (C_6H_5\ddot{N}H_3)_2PtCl_6^{-2}$ Anilinium platinic chloride Chloroplatinic acid



(2) Alkylation : Aniline reacts with alkyl halides forming secondary, tertiary and quaternary ammonium salts depending on the concentration of alkyl halides.

$$C_6H_5N(CH_3)_2 + CH_3I \longrightarrow C_6H_5(CH_3)_3NI$$

(Trimethyl phenyl ammonium iodide)

(3) **Acylation**: Aniline reacts with acid chlorides or anhydrides to form corresponding amides called anilides. [The reaction of $C_6H_5NH_9$ with benzoyl chloride is example of "Schotten Baumann reaction"]

(4) Carbylamine reaction:

$$C_6H_5NH_9 + CHCl_3 + 3KOH \longrightarrow C_6H_5NC + 3KCl + 3H_9O$$

Phenyl isocyanide (Foul smell compound)

Note: (1) Intermediate species is dichloro carbene [: CCl₂].

- (2) This is test of aniline and other primary amine, known as Isocynide test.
- (5) Hoffmann's mustard oil reaction: When aniline is heated with alc. CS₂ and excess of HgCl₂ phenyl isothiocyanate having a characteristic smell of mustard oil is formed.

$$C_6H_5NH_2 + S = C = S \xrightarrow{HgCl_2} C_6H_5N = C = S$$

Phenyl isothiocyanate

This is a test of aniline and other primary amines.

(6) Reaction with aldehydes: Aniline condenses with aldehydes to form schiff's base.

$$\begin{array}{cccc} C_6H_5NH_2 \ + \ H-C-C_6H_5 & \longrightarrow & C_6H_5N = CHC_6H_5 \ + H_2O \\ & & & & \\ O & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ &$$

(7) Reaction with Hinsberg's reagent:

$$C_6H_5 \hspace{-0.5cm} -\hspace{-0.5cm} SO_2Cl \hspace{0.1cm} + \hspace{0.1cm} HNHC_6H_5 \hspace{1cm} \hspace{0.5cm} \xrightarrow{-HCl} \hspace{0.1cm} C_6H_5SO_2NHC_6H_5$$

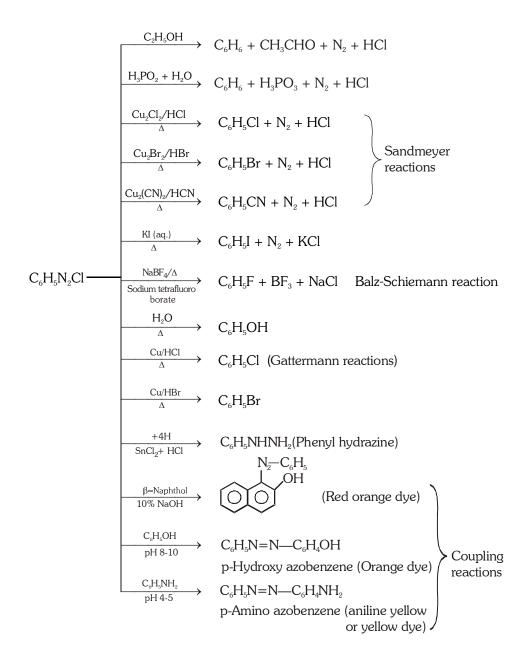
(N-Phenylbenzene sulphonamide)

(8) Diazotisation: Diazotisation is a reaction in which ice cooled solution of aniline in an inorganic acid reacts with sodium nitrite solution leading to the formation of diazonium salt.

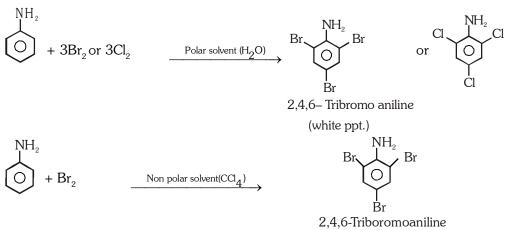
$$C_6H_5NH_2 + NaNO_2 + HCl \xrightarrow{0^{\circ}-5^{\circ}C} C_6H_5N_2Cl$$

Benzene diazonium chloride is a useful synthetic reagent. It is used in the preparation of many organic compounds





(1) **Halogenation**: In polar and nonpolar medium Chlorine and bromine react with aniline and form trichloro and tribromo aniline respectively.





Note: However, monobromo or chloro derivative of aniline can be prepared if $-NH_2$ group is first protected by acetyl group. Here the reactivity decreases due less +M effect on benzene ring.

$$\begin{array}{c}
 \text{NH}_2 \\
 \text{O}
\end{array}$$

$$\begin{array}{c}
 \text{NHCOCH}_3 \\
 \text{O}
\end{array}$$

$$\begin{array}{c}
 \text{Br}_2 \\
 \text{CH}_3\text{COOH}
\end{array}$$

$$\begin{array}{c}
 \text{NHCOCH}_3 \\
 \text{Br}
\end{array}$$

$$\begin{array}{c}
 \text{NH}_2 \\
 \text{Br}
\end{array}$$

$$\begin{array}{c}
 \text{NH}_2 \\
 \text{Br}
\end{array}$$

$$\begin{array}{c}
 \text{NH}_2 \\
 \text{Br}
\end{array}$$

o- and p-Bromo aniline

(2) Nitration:

$$\begin{array}{c}
\stackrel{\text{NH}_2}{\longrightarrow} \stackrel{\text{H}_2\text{SO}_4}{\longrightarrow} \stackrel{\text{H}_2\text{SO}_4}{\longrightarrow} \stackrel{\text{H}_2\text{SO}_4}{\longrightarrow} \stackrel{\text{NH}_2}{\longrightarrow} \stackrel{\text{NH}_2}{\longrightarrow} \stackrel{\text{NH}_2}{\longrightarrow} \stackrel{\text{NH}_2}{\longrightarrow} \stackrel{\text{NH}_2}{\longrightarrow} \stackrel{\text{NH}_2}{\longrightarrow} \stackrel{\text{NNO}_2}{\longrightarrow} \\
\stackrel{\text{NO}_2}{\longrightarrow} \stackrel{\text{NNO}_2}{\longrightarrow} \stackrel{\text{$$

(3) Sulphonation: Aniline reacts with furning H_2SO_4 to give sulphanilic acid. (p-Amino-benzene sulphonic acid.)

Sulphanilic acid

- This process is called baking.
- Sulphanilic acid is an important intermediate in the manufacturing of dyes and drugs.
- The compounds in which both proton donating & proton accepting groups present are called ampholite (dipolar ion).

(4) Catalytic hydrogenation : Aniline undergoes hydrogenation in presence of Ni at high temp. to form cyclohexanamine.

5.3 Tests of Aniline

(i) Carbylamine test: Aniline gives carbylamine test or Isocyanide test.

$$C_6H_5NH_2 + CHCl_3 + KOH \longrightarrow C_6H_5NC$$

(Bad smelling)

- (ii) Dye test: Aniline is first diazotised. On adding alkaline solution. of β -naphthol to the diazotised product a red-orange dye is formed.
- (iii) On heating with bromine water, a white ppt. is formed.



BEGINNER'S BOX-1

- **1.** Which one of following reaction is Schotten–Baumann reaction
 - (1) Acetylation of RNH₂

(2) Acylation of RNH₂

(3) Benzoylation of RNH₂

(4) All of them

- 2. Which of the following pair will yield primary amine on hydrolysis
 - (1) CH₃NCO, CH₃NC

(2) CH₃CN, CH₃NC

(3) (CH₃)₂NH, CH₃—CH=NOH

(4) None of the above

- **3.** Methylamine on treatment with chloroform and ethanolic caustic alkali gives foul smelling compound, the compound is
 - (1) CH₃NCO
- (2) CH₃CNO
- (3) CH₃CN
- (4) CH₃NC

6.0 NITRO BENZENE [C₆H₅NO₂]

It is also called as artificial oil of bitter almonds or oil of mirbane as its odour is like that of bitter almonds.

- 6.1 General Methods of Preparation
 - (1) Lab method:

$$\bigcirc + \text{ Conc. H}_2\text{SO}_4 + \text{ Conc. HNO}_3 \xrightarrow{50-60^{\circ}\text{C}}$$

(2) From diazonium salt:

$$\begin{array}{c|c} N_2Cl & NO_2 \\ \hline & & \\ \hline & &$$

6.2 Physical Properties

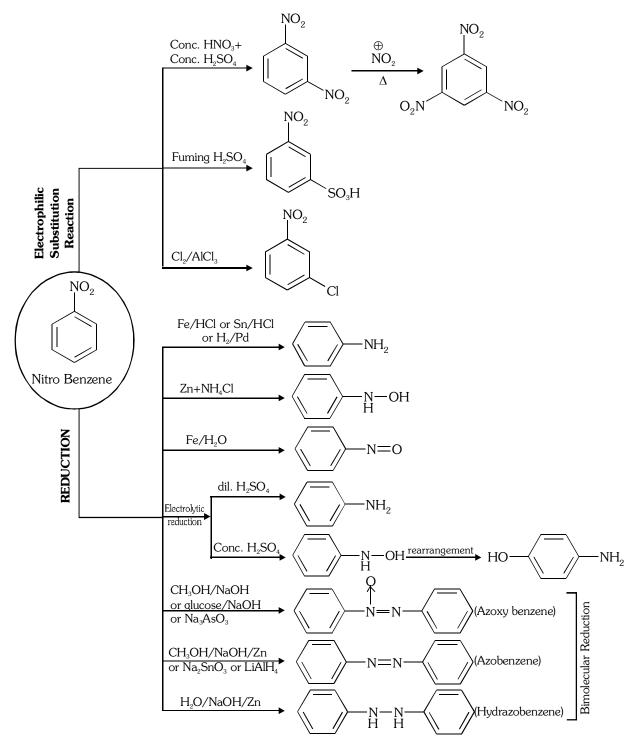
- (i) Nitro benzene is light yellow oily liquid
- (ii) It has smell of bitter almonds
- (iii) It is steam voltile. It's vapours are poisonous in nature.
- (iv) It is heavier than water
- (v) It's B. P. is 211°C
- (vi) Smell of nitro benzene and benzaldehyde is same

6.3 Chemical Properties

Nitrobezene shows following chemical reactions -

- (1) Reactions due to NO₂ group.
- (2) Reactions due to benzene ring: (A) Electrophilic substitution (B)
- (B) Nucleophilic substitution





6.4 Test of Nitrobenzene

Mullikan Barker Test: Ethanolic solution of nitrobenzene is treated with zinc dust and NH_4Cl solution. The mixture is heated and filter in a test tube containing Tollen's reagent a grey or black pricipitate (Ag mirror) is formed.

6.5 Uses

- (i) As a solvent
- (ii) In manufacture of aniline and azo dye



BEGINNER'S BOX-2

- 1. Nitration of which of the following compound is difficult :-
 - (1) Benzene
- (2) Nitrobenzene
- (3) Toluene
- (4) Phenol
- 2. Nitration of nitrobenzene in presence of fuming nitric acid will generate a :-
 - (1) Solid product
- (2) Gaseous product
- (3) Semi-solid product
- (4) Liquid product
- $\textbf{3.} \hspace{0.5cm} C_6H_5NO_2 \xrightarrow{\hspace{0.5cm} Sn/HCl} A \xrightarrow{\hspace{0.5cm} NaNO_2/HCl} B; \hspace{0.1cm} \text{To obtain benzene from } B, \hspace{0.1cm} \text{the suitable reagent is :}$
 - (1) SnCl₂ + HCl
- (2) $H_{3}PO_{2}$
- (3) C₂H₅N₂Cl
- (4) Methanol
- **4.** When nitrobenzene is heated with conc. HNO_3 and conc. H_2SO_4 the product would be obtained :-
 - (1) T.N.T.
- (2) D.N.B.
- (3) D.D.T.
- (4) T.E.L.
- 5. Which of the following has the maximum value of dipole moment?







ANSWER KEY

BEGINNER'S BOX-1	Que.	1	2	3					
	Ans.	4	1	4					
BEGINNER'S BOX-2	Que.	1	2	3	4	5			
	Ans.	2	1	2	2	2			



EXERCISE-I (Conceptual Questions)

Build Up Your Understanding

AMINE

- 1. Among the following which one is not formed in Hoffmann degradation
 - (1) RNCO
- (2) $R NH_{o}$
- (3) RCONHBr
- (4) RNC
- $CH_3CH_2CONH_2 \xrightarrow{NaOH} A,$ 2.

Aqueous solution of A

- (1) Turns blue litmus to red
- (2) Turns red litmus to blue
- (3) Does not affect the litmus
- (4) Decolourise the litmus
- 3. Ethanamine can be obtained if the following compound is heated with [KOH + Br,]
 - (1) Ethanamide
- (2) Methanamide
- (3) Propionamide
- (4) All the above
- $CH_3CONH_2 \xrightarrow{P_2O_5} A \xrightarrow{Na/EtOH} B$ 4.

Reaction II is called

- (1) Clemensen
- (2) Stephen
- (3) Mendius
- (4) Bauveault-blank reduction
- 5. CH₃CONH₂, Br₂ & KOH give CH₃NH₂ as the product. The intermediates of the reaction are :-

(a)
$$CH_3$$
— C — $NHBr$ (b) CH_3 — N = C = O

(d)
$$CH_3 - C - N < Br$$

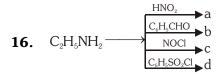
The correct answer is :-

- (1) a, b
- (2) a, c
- (3) b, d
- (4) c, d
- 6. In which case alkylamine is not formed :-
 - (1) $R X + NH_3 \longrightarrow$
 - (2) R—CH=NOH + [H] $\xrightarrow{\text{Na}}$
 - (3) R—CN + H_9O $\xrightarrow{H^+}$
 - (4) RCONH₂ + 4[H] \longrightarrow

- 7. Tertiary amine is obtained in the reaction :-
 - (1) Aniline $\xrightarrow{CH_3I}$ $\xrightarrow{CH_3I}$
 - (2) Aniline $\xrightarrow{CH_3I}$
 - (3) Nitrobenzene Sn/HCl
 - (4) None of the above
- C₂H₅NH₂ cannot be prepared by the reduction of
 - $(1) C_2H_5NO_2$
- (2) CH₃CH=NOH
- $(3) C_2H_5NC$
- (4) CH₂CN
- 9. Gabriel reaction for the synthesis of amines, involves the use of
 - (1) 1° amide
- (2) 2° amide
- (3) Imides
- (4) Aliphatic amide
- 10. Gabriel phthalimide reaction is used in the synthesis
 - (1) Primary aromatic amines
 - (2) Secondary amines
 - (3) Primary aliphatic amines
 - (4) Tertiary amines
- 11. The reaction : $[C_2H_5Br + NH_3]$ is in fact an example
 - (1) Ammonolysis only
 - (2) Nucleophilic substitution only
 - (3) Ammonolysis as well as nucleophilic substitution
 - (4) None
- **12**. Melting points are normally the highest for
 - (1) Tertiary amides
- (2) Secondary amides
- (3) Primary amides
- (4) Amines
- 13. Solubility of ethylamine in water is due to
 - (1) Low molecular weight
 - (2) Ethyl group is present in ethyl alcohol
 - (3) Formation of H-bonding with water
 - (4) Being a derivative of ammonia
- Which of the following compound liberates CO, when treated with NaHCO,
 - (1) CH₃COCH₂NH₂
- (2) CH₂NH₂
- (3) $(CH_3)_a NOH$ (4) $CH_3 NH_3 CI$



- **15.** The product obtained by the alkaline hydrolysis of C_2H_5 —N=C=O when treated with t-butyl magnesiumbromide, the compound obtained will be
 - (1) t-butylamine
- (2) n-butylamine
- (3) Isobutane
- (4) n-butane



Which product is a Schiff's base :-

- (1) a
- (2) b
- (3) c
- (4) d
- **17.** Acidic nature of amino group is shown by the reaction:
 - (1) $R-NH_2 + NOCl \rightarrow RCl + N_2 + H_2O$
 - (2) $2RNH_2 + 2Na \rightarrow 2RNH.Na + H_2$
 - (3) R.CH₂NH₂ + HNO₂ \rightarrow R.CH₂OH + N₂+ H₂O
 - (4) $R.NH_2 + HCl \rightarrow RNH_3 \stackrel{\Theta}{Cl}$
- **18.** The reagent used in the conversion of $C_2H_5NH_2$ to C_2H_5Cl would be
 - (1) SO₂Cl₂
- (2) SOCl₂
- (3) NOCl
- (4) All
- **19.** Hydrogen attached to nitrogen is released in the reaction
 - (1) RCONH₂ + NaNH₂
 - (2) $RNH_2 + Na$
 - (3) Both the above
 - (4) None of the abvoe
- **20.** If primary amines are treated with ketones the product is
 - (1) Urea
- (2) Guanidine
- (3) Amide
- (4) Schiff's base
- 21. Reactants of reaction I are

CH₃CONH₂, KOH, Br₂

Reactants of reaction-II are

CH₂NH₂, CHCl₂, KOH

The intermediate species of reaction—I and reaction—II are respectively

- (1) Carbonium ion, carbene
- (2) Nitrene, carbene
- (3) Carbene, nitrene
- (4) Carbocation, carbanion

- **22.** N, N-dimethyl acetamide is obtained in the reaction
 - (1) Acetyl chloride and methanamine
 - (2) Acetyl chloride and ethanamine
 - (3) Acetyl chloride and dimethylamine
 - (4) Acetyl chloride and diethylamine
- **23.** This compound does not respond to carbylamine reaction:-
 - (1) CH₃-CH-NH₂ CH₃
 - (2) $C_2H_5-NH-C_2H_5$
 - (3) CH₃-C-NH₂ CH₃
 - (4) CH₃-CH-CH₂-CH₃ NH₂
- **24.** Blue litmus can be turned to red by the compound (1) ROH
 - (2) RNH_o
 - (3) RNH₃ OH
 - (4) RNH₃ Cl
- **25.** Match List I with List II and select the correct answer using the codes given below:-

List I List II [Reagent] [Used as test reagent for] A. Ammonical $AgNO_3$ a. Primary amine

- B. HIO₄
- b. Aldehyde
- C Cold dil A
- o. Muerryae
- C. Cold dil. Alkaline KMnO₄
- c. Vicinal-OH groups
- D. Chloroform + NaOH d. Double bond

Cod	les :			
	Α	В	С	D
(1)	b	С	a	d
(2)	d	b	a	С
(3)	b	С	d	a
(4)	d	Ċ	b	а



- Which one of the following aromatic amino **26**. compound gives alcohol with HNO₂:-
 - (1) N,N-Dimethylaniline (2) Benzylamine
 - (3) N-methylaniline
- (4) Aniline

ANILINE

- **27**. Electrolytic reduction of nitrobenzene in weakly acidic medium gives
 - (1) Aniline
 - (2) p-Hydroxyaniline
 - (3) N-Phenylhydroxylamine
 - (4) Nitrosobenzene
- **28.** $C_6H_5NH_2 \xrightarrow{Br_2/CCl_4}$? The product is :-
 - (1) Only o- bromoaniline
 - (2) 2, 4, 6-triboromoaniline
 - (3) o-and p-bromoaniline
 - (4) Only p-bromoaniline
- 29. Before proceeding for the nitration of aminobenzene, the NH2 group is first protected by:-
 - (1) Alkylation
 - (2) Acetylation
 - (3) Formylation
 - (4) Chloromethylation
- **30.** Aniline is purified by :-
 - (1) Azeotropic distillation
 - (2) Steam distillation
 - (3) distillation in presence of magnesium
 - (4) Fractional distillation
- **31.** Reaction $C_6H_5NH_9 + HAuCl_4 \longrightarrow$

 $[C_6H_5\overset{\oplus}{N}H_3]Au\overset{\Theta}{Cl}_4$ shows ... behaviour of aniline :-

- (1) Acidic
- (2) Neutral
- (3) Basic
- (4) Amphoteric
- 32. Aniline on treatment with bromine water yields white precipitate of :-
 - (1) o-Bromoaniline
 - (2) p-Bromonailine
 - (3) 2, 4, 6-Tribromoaniline
 - (4) m-Bromoaniline

- Which compound does not show diazo reaction:-
- (3) $O_2N-\langle O \rangle -NH_2$
- Which of the following amines give N-nitroso **34**. derivative with NaNO2 and HCl:-
 - $(1) C_0H_5NH_0$

- 35. Which of the following involves nitrene as an intermediate?
 - (1) Carbylamine reaction
 - (2) Hofmann bromamide reaction
 - (3) Reimer Tiemann reaction
 - (4) Friedal crafts reaction
- **36**. Which of the following does not reduce Tollen's reagent :-
 - (1) CH₃CHO
- (2) HCOOH
- (3) C₆H₅NHOH
- $(4) C_6 H_5 N H_9$
- **37.** Aniline can be obtained by :-
 - (1) Benzoyl chloride and ammonia
 - (2) Reduction of benzamide
 - (3) Phenol and ammonia in presence of ZnCl₂
 - (4) Benzoic anhydride and ammonia
- Aniline on direct nitration produces :-
 - (1) o-Nitroaniline
- (2) m-Nitroaniline
- (3) p-Nitroaniline
- (4) All
- **39**. Nitration of acetanilide followed by hydrolysis gives
 - (1) o-Nitroaniline
- (2) p-Nitroaniline
- (3) o- & p-Nitroaniline (4) o-Nitroanilinium ion
- **40.** $C_6H_5NH_2 \xrightarrow{NaNO_2/HCl} A$, Which is the incorrect structure of the product 'A' :-
 - (1) C_6H_5 —N=N—Cl (2) $[C_6H_5N_2]$ Cl
 - (3) $[C_{\epsilon}H_{\epsilon} N \equiv N] \stackrel{\Theta}{C}$ (4) $[C_{\epsilon}H_{\epsilon} N \equiv N] \stackrel{\Theta}{C}$



- **41.** Chloroform and ethanolic KOH is used as a reagent in the following reaction :-
 - (a) Hoffmann carbylamine reaction
 - (b) Hoffmann degradation reaction
 - (c) Reimer-Tiemann reaction
 - (d) Hoffmann mustard oil reaction

Code is :-

- (1) Only for a
- (2) Only for a and b
- (3) Only for b and d
- (4) Only for a and c
- **42.** Acetanilide when treated with bromine in acetic acid mainly gives :-
 - (1) o-Bromoacetanilide
 - (2) N-Bromoacetanilide
 - (3) p-Bromoacetanilide
 - (4) m-Bromoacetanilide
- **43.** Aromatic nitriles (ArCN) are not prepared by reaction
 - (1) ArX + KCN
 - (2) $ArN_2^+ + CuCN$
 - (3) ArCONH₂ + P_2O_5
 - (4) ArCONH₂ + SOCl₂

NITRO GROUP, CYANIDE & ISOCYANIDE

44. Aniline in a set of reactions yielded end product D

$$C \xrightarrow{HNO_2} D$$

The structure of the product D would be

- (1) $C_6H_5CH_9OH$
- (2) $C_6H_5CH_2NH_2$
- (3) C₆H₅NHOH
- (4) C₆H₅NHCH₂CH₂
- **45.** In the reaction sequence identify the functional group present in A, B, C:-

$$A \xrightarrow{Sn/HCl} B \xrightarrow{HNO_2} C \xrightarrow{C_2H_5OH} C_6H_6$$

- (1) NO₂, NH₂, N=N
- (2) NO₂, NH₂, OH
- (3) —OH, —NH₂,—NO
- (4) $-NH_2$, $-NO_2$, -N=N-

46. ϕ — $X \xrightarrow{NaNO_2/HCl} C_6H_5N_2Cl \xrightarrow{Water} \phi$ —Y,

In the above sequence X and Y are :-

- (1) o-, p- and m-directing
- (2) o-, p- and o-, p-directing
- (3) m and m directing
- (4) m and o, p directing
- **47.** Which of the following compound gives an explosive on decarboxylation:-
 - (1) 2,4, 6-Trinitrobenzoicacid
 - (2) 2, 4-Dinitrobenzoicacid
 - (3) o-Aminobenzoicacid
 - (4) o-Hydroxybenzoicacid
- **48.** The gas leaked from a storage tank of the Union Carbide plant in Bhopal gas tragedy was:-
 - (1) Methylisocyanate
- (2) Methylamine
- (3) Ammonia
- (4) Phosgene

49.
$$CH_3$$
— CH_3 — CH_3 reduction ?

- (4) None
- **50.** Reaction of RCN with sodium and alcohol leads to the formation of :-
 - (1) RCONH_o
- (2) RCOO⁻NH₄⁺
- (3) RCH₂NH₂
- (4) R(CH₂)₂NH₂



- **51.** $C_6H_5NO_2 \xrightarrow{SnCl_2/HCl} A \xrightarrow{NaNO_2/HCl} B$; In the above sequence Benzene from B, is suitably obtained by using :-
 - (1) Ethanol
 - (2) $H_{3}PO_{2}$
 - (3) Both the above
 - (4) Methanol
- **52**. Which reagent is used to get iodo benzene from benzene diazonium acid sulphate $[C_6H_5N_9HSO_4]$:
 - (1) CuBr, ∆
- (2) Cu powder + HI
- (3) KI, Δ
- (4) None
- **53.** Which of the following is used as a solvent in the Friedel-Crafts reaction :-
 - (1) Toluene
- (2) Nitrobenzene
- (3) Benzene
- (4) Aniline

Match list I with II and choose the correct answer from the codes given below:-

List I

- (A) Aniline
- a. Used in making azo
 - dyes
- (B) Nitrobenzene
- b. Sulpha drug
- (C) Sulphanilamide c. Solvent in the Friedel
 - Crafts reaction
- (D) Trinitrotoluene d. Used as explosive

Code is :-

	Α	В	C	D
(1)	a	С	b	d
(2)	a	b	С	d
(3)	С	d	a	b
(4)	d	C	b	а

- **55.** In the Sandmeyer's reaction, -N=N-X group of diazonium salt is replaced by :-
 - (1) Halide group
- (2) Nitro group
- (3) —OH group
- (4) —NHNH₂ group

EX	EXERCISE-I (Conceptual Questions)								ANSWER KEY						
Que.	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
Ans.	4	2	3	3	1	3	1	3	3	3	3	3	3	4	3
Que.	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30
Ans.	2	2	3	3	4	2	3	2	4	3	2	1	2	2	2
Que.	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45
Ans.	3	3	4	3	2	4	3	4	3	1	4	3	1	1	1
Que.	46	47	48	49	50	51	52	53	54	55					
Ans.	2	1	1	2	3	3	3	2	1	1					



Directions for Assertion & Reason questions

These questions consist of two statements each, printed as Assertion and Reason. While answering these Questions you are required to choose any one of the following four responses.

- (A) If both Assertion & Reason are True & the Reason is a correct explanation of the Assertion.
- **(B)** If both Assertion & Reason are True but Reason is not a correct explanation of the Assertion.
- **(C)** If Assertion is True but the Reason is False.
- **(D)** If both Assertion & Reason are false.
- **1. Assertion**: Boiling point of trimethyl amine is higher than that of n-propyl amine.

Reason: H-bonding is stronger in tertiary amines.

- (1) A
- (2) B
- (3) C
- (4) D
- 2. Assertion :- $H-C-N-CH_3$ and $\parallel \ \mid \ O \ CH_3$

$$CH_3-C-NH-CH_3$$
 are metamers 0

Reason :- There is adjustment of $-CH_2$ - group in the molecule.

- (1) A
- (2) B
- (3) C
- (4) D
- **3. Assertion**: Pyrole is alicyclic compound.

Reason: It is aliphatic in nature and has cyclic chain.

- (1) A
- (2) B
- (3) C
- (4) D
- **4. Assertion:** The following compound are given below –

$$\begin{array}{c} \mathrm{CH_{3}CH_{2}NO_{2}\,,CH_{3}-\!CH.NO_{2}} \,\,\& \\ \mathrm{CH_{3}} \end{array}$$

$$\begin{array}{c} CH_3-CH-CH_2-CH_3 \\ | \\ NO_2 \end{array}$$

all three compounds are soluble in NaOH.

Reason :- All above compound have lower boiling than $\mathrm{CH_3NO_2}$.

- (1) A
- (2) B
- (3) C
- (4) D

5. Assertion: Acetonitrile is another name of ethanenitrile.

Reason :- α -H atom of acetonitrile exhibit acidic character.

- (1) A
- (2) B
- (3) C
- (4) D
- **6. Assertion**: Benzene diazonium chloride on reaction with ethanol gives benzene.

Reason:- Ethanol is good oxidising agent.

- (1) A
- (2) B
- (3) C
- (4) D
- **7. Assertion**: Diazonium salt is prepared at low temperature.

Reason: Diazonium salt is hydrolysed to give phenol.

- (1) A
- (2) B
- (3) C
- (4) D
- **8. Assertion**: Carbylamine reaction involves reaction between 1° amine and chloroform in basic medium to give isocyanide.

Reason: Methylene (: CH_2) carbene is formed as a reaction intermadiate.

- (1) A
- (2) B
- (3) C
- (4) D
- Assertion: Suitable method for preparation of primary amine is when haloalkane is treated with excess of NH₃.

Reason: In this reaction primary, secondary, tertiary amines are formed.

- (1) A
- (2) B
- (3) C
- (4) D

ANSWER KEY

Que.	1	2	3	4	5	6	7	8	9	
Ans.	4	4	4	3	2	3	1	3	3	